## **Summary of Quality Control Samples and the Information They Provide**

Data Quality Indicator	QC Check and QC Sample			Purpose								
		Sample Collection				Sample Transport	L	aboratory/l	od	To evaluate or determine the source of		
		Sampling Equipment	Conditions During Sampling	Preservation Technique	Sampling Matrix	Shipment Process	Sample Storage at Laboratory	Sample Preparation Reagents	Sample Preparation Equipment	Analytical Methods Reagents/ Standards	Analytical Equipment	measurement error arising from:
Accuracy/Bias (Positive bias introduced by contamination)	Equipment Blank (Rinsate Blank)	11		1		(VOCs)	✓	11	11	11	11	Carryover contamination resulting from successive use of sampling equipment. Also see preparation blank.
	Pour Blank (Ambient, or Field Blank)		11	1		(VOCs)	✓	11	11	11	11	Ambient contamination of sample during sampling exercise (e.g., sand blowing into a water-metals sample, or high concentration of volatiles in air). Also see preparation blank.
	Volatile Organic, or Radiological (Radon) Trip Blank					11	11	11	11	11	11	Contamination introduced during shipment. Usually limited to VOCs and radiological parameters such as radon. Also see preparation blank.
	Volatile Organic Storage Blank (Refrigerator blank)						11			✓	1	Cross contamination introduced during sample storage usually for VOCs. Also may be used for radon, tritium.
	Reagent Blank (one per lot number)			11				1	1	1	1	Contamination introduced by reagents used as sample preservatives.
	Preparation Blank							11	11	11	11	Contamination introduced by preparation process, glassware, analytical reagents, and analytical instrumentation.
	Instrument (System) Blank										11	Contamination originating with the analytical equipment.
Accuracy/Bias (Bias due to inadequate temperature control)	Cooler Temperature Blank			11		11						Inadequate temperature control during shipping leading to potential loss of target analytes.

<sup>✓✓ =</sup> Primary purpose of QC sample ✓ = Secondary purpose of QC sample

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Indicator			Sample 0	Collection		Sample Transport	L	aboratory/l	od	To evaluate or determine the source of		
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Accuracy/Bias (Bias due to sample matrix or sample preparation/ analytical	Matrix Spike				11			1	1	✓	1	Preparatory and analytical bias for specific compounds in specific sample matrices.
	Surrogate Spike				11			1	1	1	1	Preparatory and analytical bias in specific sample matrices.
methodology/ operator error)	Laboratory Control Samples							11	11	11	11	Laboratory's ability to accurately identify and quantitate target compounds in a reference matrix at a known concentration.
	Performance Evaluation Sample- Ampule Single Blind							11	11	11	11	Laboratory's ability to accurately identify and quantitate target compounds.
	Performance Evaluation Full Volume-Double Blind Prepared in Site-specific Matrix				11		11	11	11	11	11	Laboratory's ability to accurately identify and quantitate target compounds in a reference matrix.
	Laboratory Fortified Blank							11	11	11	11	Method preparatory and analytical sensitivity and bias for specific compounds in a reference matrix at the quantitation limit.
	Initial Calibration									11	11	Sets the response to a known concentration to ensure the instrument will produce acceptable quantitative data.
	Continuing Calibration & Verification									<b>//</b>	11	Checks the accuracy and stability of the instrument response.
	Instrument Performance Check Sample									11	11	Checks that an instrument can accurately identify and quantitate target analytes at specific concentration levels.

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Accuracy/Bias (Bias due to methodology)	Field Splits (Homogenized Samples)							<b>//</b>	11	11	11	Comparability of results between two methods, or laboratories (e.g., field and fixed).
	Field Splits (extracts)							11	11	11	11	Comparability of results between two methods, or laboratories when sample matrix is known to be extremely heterogenous.
Sensitivity	Laboratory Fortified Blank							11	11	11	11	Laboratory preparatory and analytical sensitivity and bias for specific compounds in a reference matrix at quantitation limit concentrations.
	Method Detection Limit Studies				11			11	11	11	11	Statistical determination that defines the minimum concentration of a substance that can be measured and reported with 99% confidence that the analyte concentration is > zero.
	Low Point of Initial Calibration Curve									11	11	Instrument is capable of producing acceptable qualitative and quantitative data at the lowest concentration that sample results will be reported; the quantitation limit.

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Precision	Field Duplicates	✓	1	1	11	(VOCs)	✓	1	1	✓	1	Cumulative effects of both field and laboratory precision to measure overall precision.
	Laboratory Duplicates				11			1	1	✓	1	Laboratory preparatory and analytical precision.
	Matrix Spike Duplicates				11			1	1	1	1	Laboratory and analytical bias and precision for specific compounds in specific sample matrices.
	Analytical Replicates										11	Analytical precision for determinative instrumentation.
	Internal Standards										<b>//</b>	Instrument precision and stability.
Representa- tiveness	Field Replicates (not homogenized), also known as co- located samples	1	1	1	11	(VOCs)	1	1	1	<b>√</b>	1	Comparability of samples taken closely in time or space.

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